This Page Is Inserted by IFW Operations and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

As rescanning documents will not correct images, please do not report the images to the Image Problem Mailbox.

÷			
<u>;:</u>			
·			
		•	
			,



WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6: B32B 27/18, B65D 81/26, A23L 3/34

(11) International Publication Number:

WO 97/32722

(43) International Publication Date: 12 September 1997 (12.09.97)

(21) International Application Number:

PCT/US97/03307

A1

(22) International Filing Date:

26 February 1997 (26.02.97)

(30) Priority Data:

08/612.510

7 March 1996 (07.03.96)

US

(71) Applicant: CHEVRON CHEMICAL COMPANY [US/US]; P.O. Box 5047, San Ramon, CA 94583-0947 (US).

(72) Inventors: CHING, Ta, Yen; 10 Santa Yorma Court, Novato, CA 94945 (US). GOODRICH, Joseph, L.; 3545 Wilkinson Lane, Lafayette, CA 94549 (US). KATSUMOTO, Kiyoshi; 2615 Brooks Avenue, El Cerrito, CA 94530 (US).

(74) Agents: MICHEL, Marianne, H. et al.; Chevron Corporation. Law Dept., P.O. Box 7141, San Francisco, CA 94120-7141 (US).

(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).

Published

With international search report.

Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.

(54) Title: OXYGEN SCAVENGING SYSTEM INCLUDING A BY-PRODUCT NEUTRALIZING MATERIAL

(57) Abstract

Provided is a system useful for oxygen scavenging which comprises at least two components, i.e., an oxygen scavenging material which forms at least one by-product upon reaction thereof with oxygen, and an effective amount of a neutralizing material capable of neutralizing at least a portion of theses by-products. In a preferred application, a multi-layer structure that can be employed in producing packages and in particular food packages, comprises a first layer including an oxygen scavenging material and a second layer which includes a material that is capable of neutralizing at least a portion of the by-products produced by the oxidation of the oxygen scavenging material within the first layer. These two layers are arranged such that, upon formation of the package, the second layer is interior to the first layer. Furthermore, the multi-layer film can include one or more of an oxygen barrier layer, a polymeric selective barrier layer, and a heat-sealable layer.

> Serial No. 10/717,297 ` Ref. B4

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

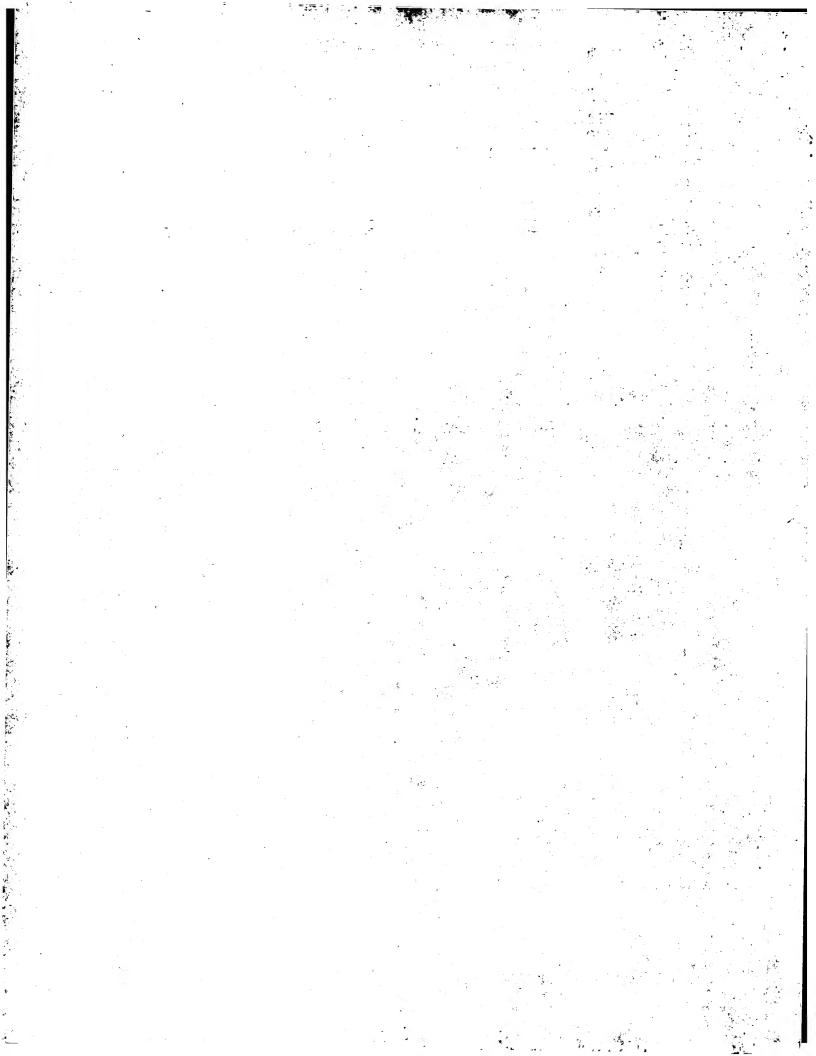
AM	Armenia	GB	United Kingdom	MW	Malawi
AT	Austria	GE	Georgia	MX	Mexico
AU	Australia	GN	Guinea	NE	Niger
BB	Barbados	GR	Greece	NL	Netherlands
BE	Belgium	HU	Hungary	NO	Norway
BF	Burkina Faso	IE	Ireland	NZ	New Zealand
BG	Bulgaria	IT	Italy	PL	Poland
BJ	Benin	JP	Japan	PT	Portugal
BR	Brazil	KE	Kenya	RO	Romania
BY	Belarus	KG	Kyrgystan .	RU	Russian Federation
CA	Canada	KP	Democratic People's Republic	SD	Sudan
CF	Central African Republic		of Korea	SE	Sweden
CG	Congo	KR	Republic of Korea	SG	Singapore
CH	Switzerland	KZ.	Kazakhstan	SI	Slovenia
CI	Côte d'Ivoire	LI	Liechtenstein	SK	Slovakia
CM	Cameroon	LK	Sri Lanka	SN	Senegal
CN	China	LR	Liberia	SZ	Swaziland
CS	Czechoslovakia	LT	Lithuania	TD	Chad
CZ	Czech Republic	LU	Luxembourg	TG	
DE	Germany	LV	Latvia	TJ	Togo Tajikistan
DK	Denmark	MC	Monaco	TT	•
EE	Estonia	MD	Republic of Moldova	UA	Trinidad and Tobago Ukraine
ES	Spain	MG	Madagascar	UG	
FI	Finland	ML	Mali	US	Uganda
FR	France	MN	Mongolia	UZ	United States of America
GA	Gabon	MR	Mauritania		Uzbekistan .
	· ·	172.00	AT ASSESS ASSESSED.	VN	Viet Nam

INTEL ATIONAL SEARCH REPORT

information on patent family members

Inter mal Application No
PCT/US 97/03307

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0638487 A	15-02-95	JP 7069380 A	14-03-95
WO 9301049 A	21-01-93	AU 2307292 A EP 0526977 A	11-02-93 10-02-93
US 5211875 A	18-05-93	AU 650140 B AU 1811192 A AU 659773 B AU 5930594 A CA 2071079 A EP 0520257 A HU 66841 A,B IL 102160 A JP 5194949 A SK 197192 A US 5425896 A US 5498364 A	09-06-94 07-01-93 25-05-95 16-06-94 28-12-92 30-12-92 30-01-95 27-11-95 03-08-93 07-09-94 20-06-95 12-03-96
EP 0464782 A	08-01-92	AT 143857 T AU 638845 B AU 7946991 A CA 2043301 A,C DE 69122553 T ES 2094774 T JP 4232740 A JP 7002405 B NO 176789 B US 5290613 A	15-10-96 08-07-93 09-01-92 04-01-92 14-11-96 15-05-97 01-02-97 21-08-92 18-01-95 20-02-95 01-03-94
EP 0720821 A	10-07-96	AU 4062695 A CA 2165709 A CN 1132608 A JP 8229388 A	04-07-96 28-06-96 09-10-96 10-09-96



-1-

1	Catgen Scavenging Ststem including
2	a by-product neutralizing material
3	
4	BACKGROUND OF THE INVENTION
5	
6	The present invention relates to an improved oxygen scavenging system for use
7	with oxygen sensitive materials, e.g., food, which system finds particular utility in
8	the multi-layer films that includes an oxygen scavenging layer.
9	
10	Oxygen scavenging materials have been developed partly in response to the
11	food industry's needs of having longer shelf-life and better flavor preservation fo
12	packaged food.
13	
14	These oxygen scavenging materials constitute at least a portion of the food
15	package, and these materials remove oxygen which surrounds the food product,
16	e.g., the residue oxygen in a package after sealing or air which has leaked into
17	the package, thereby inhibiting spoilage of the food.
18	·
19	One method which is currently being employed involves the use of "active
20	packaging" where the food product package is modified in some way to control
21	the exposure of the food product to oxygen. Such "active packaging" can
22	include sachets containing compositions, e.g., iron-based compositions such as
23	Ageless® which scavenges oxygen in the package through an oxidation
24	reaction. However, sachets are not advantageous for a variety of reasons not
25	the least of which is the problems caused by the accidental ingestion of the
26	sachets or the material present therein.
27	

Recent attempts have involved incorporating an oxygen scavenger into the 1 package structure itself. In such an arrangement, oxygen scavenging materials 2 constitute at least a portion of the food package. One example of such an 3 arrangement comprises a scavenging wall which includes inorganic powders 4 and/or salts. See, for example, European Applications 367,835; 366,245; 5 367,390; and 370,802. However, such composition is difficult, if not impossible, 6 to adequately process and thereby is not considered commercially feasible. 7 8 In another more promising arrangement, oxygen scavenging materials can be 9 low molecular-weight oligomers that are typically incorporated into processable 10 11 packaging polymers or can be oxygen scavenging polymers in which either the backbone is designed to break apart when the polymer reacts with oxygen or in 12 13 which, initially at least, pendant oxidizable side chains react with oxygen. 14 Other methods which have been disclosed include that of European Patent 15 Application 0454437, wherein an oxygen absorbent composition is disclosed. 16 The composition contains a linear hydrocarbon polymer having one or more 17 unsaturated groups or a mixture of linear hydrocarbon polymers having one or 18 19 more unsaturated groups with an unsaturated fatty acid compound and an oxidation promoter as essential components. And, optionally, the composition 20 21 can contain a basic substance and/or an adsorption substance. 22 23 European Patent Application 0424855 discloses an inhibitor parcel comprising a composition containing an unsaturated fatty acid compound as its main 24 ingredient and a permeable diffusing parceling material prepared by laminating 25 26 and bonding an oxygen permeable resin layer onto one side of a base sheet 27 made of a fibrous material and an adhesive, and then laminating and bonding thereunto an oxygen permeable resin layer, and then laminating and bonding a 28

porous film of a low softening point resin or a low softening point unwoven fabric 1 onto the other side of the base sheet. The parcel is directed to preserving 2 electronic devices and electronic parts and excluding oxygen as well as water. 3 4 Systems have also been designed to control odors which can be generated by 5 deterioration of the contents (food) being stored, or deterioration of the film 6 packaging used in storing the contents. For example, in Japanese Kokai Patent 7 No. HEI6-223925, published February 1, 1994, a deodorant packaging film is 8 described. The packaging film is obtained by constructing a film having inner 9 and outer surface layers of polypropylene resin that contains a polybutene or 10 polyisobutylene, tackifier, and deodorant and constructing the middle layer from 11 polyethylene terephthalate or a polyamide resin. The deodorant component of 12 the polypropylene resin composition is described as being flavonoid deodorants, 13 polyphenol components containing deodorants, deodorants that have turpentine 14 oil as a major component, and deodorants based on organic acids. 15 16 U.S. Patent No. 5,340,884 discloses a polyester/polyamide blend having an 17 excellent gas barrier property and an improved flavor retaining property. In 18 particular, the polyethylene terephthalate/low molecular weight-polyamide blend 19 reduces the concentration of acid aldehyde contained in the polyester, thereby 20 21 rendering the blend more suitable for the storage of food. 22 U.S. Patent No. 5,284,892 describes a system which is an aldehyde scavenging 23 composition. Such compositions are useful in producing packaging films for oil 24 containing foods, which give off aldehydes. The compositions comprises a 25 polyalkylene imine and a polyolefin polymer. See also U.S. Patent 26 27 No. 5,362,784. 28

/32722 PCT/US97/03307

International Publication No. WO 93/01049 discloses a packaging composition 1 that helps to extend shelf life of oil containing foods by absorbing undesirable 2 aldehydes that are produced during oil degradation. The composition suggested 3 to be used in absorbing the aldehydes are primary and/or secondary amine 4 groups and strong inorganic bases. 5 6 7 European Patent Application 0504726 discloses a food preserving agent which comprises an oxygen absorbent and a substance which is able to remove 8 9 acetaldehyde. 10 European Patent Application 0464782 is directed to a multi-layer thermoplastic 11 12 film having a vinylidine chloride polymer barrier and at least a polyolefin second 13 layer laminated to the barrier. Upon irradiation of the barrier layer, odor is 14 generated due to ionizing irradiation of the barrier layer. Hydrotalcite is blended with the polyolefin in order to substantially reduce the odor generated by the 15 vinylidine chloride copolymer barrier. 16 17 Japanese Kokai HEI5-247276 discloses an oxygen barrier resin composition. 18 19 The oxygen barrier resin composition comprises a polyolefin, an oxidation catalyst and an odor absorbent. The odor absorbent is used to suppress the 20 21 odor caused by oxidation of the polyolefin. As suitable absorbents, there are 22 mentioned natural zeolite, synthetic zeolite, silica gel, activated carbon, 23 activated clay, activated aluminum oxide, magnesium silicate, aluminum silicate 24 and the like. Synthetic hydrotalcite type compounds can also be used. 25 A major problem that still needs to be addressed, however, is that a wide variety 26 27 of organic compounds are produced upon oxidation of the oxygen scavenging material. Many of these oxidation products can migrate from the layer carrying 28

•	an oxygen seavenging material and enter the air surrounding the food or even
2	enter the food itself.
3	
4	Such oxidation products can have foul odors or can even be compounds that are
5	undesirable. It is therefore highly desirable to provide a way to prevent such
6	oxidation products from entering a packaged volume that contains food.
7	
8	One attempt to solve the problem of migration of oxidation products involves the
9	use of a composition comprising two layers, where one layer carries an oxygen
10	scavenging material and one layer is a barrier situated between the packaged
11	volume and the layer carrying an oxygen scavenging material. See U.S.
12	Application Serial No. 08/304,303, filed September 12, 1994, which is
13	incorporated herein by reference for all purposes.
14	
15	One problem with this approach is that many barriers are capable of blocking the
16	migration of molecules of certain sizes, but they fail in blocking very small
17	organic molecules. On the other hand, there are good barriers which are
18	effective to block the migration of small oxidation products into, e.g., the
19	enclosed volume of the package, but they also slow the oxygen migrating from
20	the enclosed volume to the organic scavenging material.
21	
22	Other multi-layer film packaging materials are also recognized within the art.
23	Such films can include, in addition to the layers discussed above, a "polymeric
24	selective barrier layer' such as that described in U.S. Application Serial
25	No. 08/304,303, filed September 12, 1994. While such selective barrier layers
26	can prevent the migration of certain, in fact many, of the undesirable oxidation
27	products, certain by-products, and in particular, those having a smaller

molecular size may not be effectively blocked. This is particularly true for those 1 2 by-products which are of a similar or slightly larger size to gaseous oxygen. 3 Thus, the need still exists for an improved system for protecting oxygen sensitive 4 materials, and in particular a system applicable to multi-layer film for use in food 5 packaging which is capable of neutralizing such molecules, thereby reducing 6 those problems associated with these by-products as discussed above. 7 8 9 SUMMARY OF THE INVENTION 10 11 Among other aspects, this invention is based upon the surprising discovery that an oxygen scavenging system, particularly one useful in films, can be improved 12 through the use of oxidation by-product neutralizing materials in a separate 13 14 layer. By a "neutralizing" material is meant a material which can react with, complex with and overall eliminate the troublesome oxidation by-product 15 compound. Where the by-product is an acid, it can be neutralized as in an 16 17 acid/base reaction by the neutralizing material. Where the by-product is not acid, the compound can be neutralized by other understood chemical 18 interactions with the neutralizing material of the present invention. Therefore, 19 the term "neutralize", in the context of the present invention, is intended to be 20 21 broader than the literal acid/base reaction. 22 In one aspect, the present invention relates to a multi-layer structure comprising: 23 24 25 a first layer which includes an oxygen-scavenging material which react with 26 oxygen by way of an oxidation reaction; and

PCT/US97/03307

1	a second layer including an effective amount of a neutralizing material capable
2	of neutralizing at least a portion of the by-products of the oxidation reaction.
3	Moreover, these two layers are preferably arranged such that the second layer is
4	located between the first layer and the interior of the package.
5	
6	Moreover, this multi-layer structure preferably further includes an oxygen barrier
7	layer located outside of the first layer, and/or a polymeric selected barrier layer
8	which is also located inside the first layer.
9	
10	The oxygen scavenging system according to the present invention is preferably
11	employed in a packaging environment which packaging holds an oxygen
12	sensitive material. Such materials can include food, cosmetic/beauty materials
13	and other chemicals, as well as electronic materials.
14	
15	In the present invention, the oxygen scavenging material is preferably an organic
16	oxygen scavenging material while the neutralizing material is preferably selected
17	from the group consisting of inorganic bases and organic bases.
18	·
19	BRIEF DESCRIPTION OF THE DRAWING
20	
21	The figure of the Drawing illustrates one embodiment of a multi-layer film
22	according to the present invention.
23	
24	DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS
25	
26	As discussed above, the oxygen scavenging system of the present invention
27	includes two components contained in separate layers, i.e., an oxygen

scavenging material, and an oxidation by-product neutralizing material. Each of 1 2 these two components will now be discussed in more detail. 3 The first component, the oxygen scavenging material, may be any oxygen 4 scavenging material recognized in the art. Suitable examples of such materials 5 include unsaturated organic compounds such as carotene, ascorbic acid, 6 squalene, or dehydrated caster oil. Suitable materials are also disclosed in 7 EP 0 507 207, which is incorporated by reference in its entirety herein. 8 9 The oxygen scavenging material is maintained in layers separate from the 10 11 neutralizing material layer. This is important, as it has been found that a mixing of the two components can cause deleterious effects on the oxygen scavenging 12 13 properties of the oxygen scavenger, thereby defeating the entire purpose of the 14 system. The two components are thereby maintained in different layers, and 15 surprisingly offers excellent oxygen scavenging properties as well as protection for the packaged material from the oxidation by-products. 16 17 This scavenging compound may be introduced in a variety of manners, 18 19 depending on the ultimate use of the system. For example, it may be blended 20 with a carrier, e.g., polymer, which itself may or may not scavenge oxygen, or it 21 may be coated onto a material such as aluminum foil or paper or even be incorporated into a material such as paper. The oxygen scavenging material 22 may be in localized areas on a layer, e.g., the oxygen scavenging material may 23 24 be in a patch that is laminated to another layer in a multi-layer arrangement. 25 The oxygen scavenging material can further be a polymer having oxidizable 26 27 sites in the polymer and containing a catalyst such as a transition metal salt that

7	assists initiation of oxidation of the oxidizable sites. Such a material is
2	particularly useful when the scavenging material forms a layer of a film.
3	
4	Examples of polymers having oxidizable sites include polybutadiene or other
5	polymers containing unsaturated sites, such as disclosed in U.S. Pat.
6	No. 5,211,875; poly(meta-xylenediamine-adipic acid) (also known as MXD-6),
7	disclosed in U.S. Pat. Nos. 5,021,515 and 5,049,624 and EP 0 519 616; and
8	poly(ethylene- methyl acrylate-benzyl acrylate), disclosed in U.S. Ser.
9	No. 08/091,120, filed August 12, 1993, inventors T. Ching, K. Katsumoto,
10	S. Current and L. Theard, each of which is incorporated by reference in its
11	entirety herein.
12	•
13	For sake of completeness, poly(ethylene-alkyl acrylate-benzyl acrylate) can be
14	made, e.g., by solution transesterification. An ethylene-alkyl acrylate copolymer
15	such as ethylene-methyl acrylate copolymer is dissolved in an appropriate
16	solvent such as decalin, and heated to and maintained at reflux in the presence
17	of an effective amount of a transesterification catalyst, such as tetraethyl titanate
18	or di-butyl tin laureate, and an alcohol containing a benzyl radical, such as
19	benzyl alcohol. The solution is then cooled, and the polymer is precipitated in
20	methanol and dried in a vacuum oven.
21	
22	An effective amount of a transition metal salt catalyst such as cobalt
23	neodecanoate is incorporated into the precipitated polymer by melting the
24	polymer in, for example, an extruder, and mixing the salt dissolved in a solvent
25	such as hexane into the polymer melt.
26	
27	The transesterification above may also occur using a melted ethylene-alkyl
28	acrylate copolymer in a reactive extruder maintained at transesterification

conditions and in the presence of an effective amount of a transesterification 1 2 catalyst and an alcohol containing a benzyl radical. The above-described process can be used to transesterify ethylene alkyl acrylate copolymers to yield 3 compounds suitable for use as neutralizing amine-containing polymers in this 4 5 invention. 6 The transition-metal salt that assists initiation or catalyzes the oxidation of the 7 8 oxidizable sites generally comprises an element chosen from the first, second and third transition series of the periodic table of the elements. This transition-9 metal salt is in a form which facilitates or imparts scavenging of oxygen by the 10 afore-described polymers. It is generally believed that the transition-metal salt is 11 in an ionic state such that the transition element can readily inter-convert 12 13 between at least two oxidation states. 14 Suitable transition-metal elements include, but are not limited to, manganese II 15 or III, iron II or III, cobalt II or III, nickel II or III, copper I or II, rhodium II, III or IV, 16 and ruthenium. The oxidation state of the transition-metal element when 17 18 introduced into the composition is not necessarily that of the active form. It is only necessary to have the transition-metal element in its active form at or 19 shortly before the time that the composition is required to scavenge oxygen. 20 21 22 The transition-metal element is preferably iron, nickel or copper, more preferably 23 manganese and most preferably cobalt. 24 25 Suitable counter-ions for the transition metal element are organic or inorganic anions. These include, but are not limited to, chloride, acetate, oleate, linoleate, 26 caprylate, stearate, palmitate, 2-ethylhexanoate, citrate, glycolate, benzoate, 27 28 neodecanoate or naphthenate. Organic anions are preferred.

1	
2	Preferable salts include cobalt (II) 2-ethylhexanoate, and cobalt benzoate. More
3	preferable salts include cobalt (II) neodecanoate, cobalt (II) oleate, cobalt (II)
4	linoleate, and cobalt (II) caprylate.
5	•
6	The transition-metal element may also be introduced as an ionomer, in which
7	case a polymeric counter-ion is employed. Such ionomers are well known in the
8	art. See U. S. Patent No. 3,264,272, which is incorporated by reference in its
9	entirety.
10	
11	The oxygen scavenging material comprised of a polymer and transition metal
12	salt contains a sufficient quantity of the transition-metal salt to promote oxygen
13	scavenging in the polymer. Generally, this requires a ratio of moles to benzyl
14	radicals to moles of transition-metal element between about 2000:1 to about
15	10:1. Preferably, this molar ratio is between 200:1 and 20:1. The preferred
16	amount of transition-metal element will typically vary with which transition-metal
17	salt is used.
18	
19	As discussed above, the oxygen scavenging material may be introduced into the
20	system by any of a variety of techniques. For example, in forming the multi-layer
21	structure, it can be blended into a composition for forming a layer of the structure
22	or laminated or sprayed onto the formed layer, and/or may be a layer itself.
23	
24	In particular, the oxygen scavenging material may be coated onto a polymer
25	layer or onto a multi-layer structure, in which case the oxygen scavenging
26	material normally forms its own layer. The particular method of introduction
27	selected is dependent upon the particular scavenging material employed.
28	

1	The oxygen scavenging material is preferably present in an amount sufficient to
2	scavenge at least 0.1 cc O ₂ /gram of oxygen scavenging material/day.
3	Preferably, it is capable of scavenging at least about 0.5, and more preferably at
4	least about 1 cc O ₂ /gram of oxygen scavenging material/day.
-5	The state of the s
6	As discussed above, the oxygen scavenging material, upon consumption of
7	oxygen, typically produces certain by-products, e.g., volatile organic oxidation
8	products. Examples of these oxidation by-products include carboxylic acids,
9	such as acetic, propionic, butyric, valeric and benzoic acids; aldehydes, such as
10	heptanal and benzaldehyde; ketones, such as acetone and methyl ethyl ketone;
11	esters, such as methyl formate; alcohols, and the like. These by-products can
12	cause problems.
13	
14	However, the second component present in the oxygen scavenging system is a
15	material capable of neutralizing at least a portion of the oxidation by-products
16	produced from the reaction of oxygen with the oxygen scavenging material.
17	Generally, the neutralizing material in the second layer comprises an acid-,
18	alconor- or aldehyde-reactive material. The particular material selected is
19	dependent upon the scavenging material employed and the by-products whose
20	migration needs to be controlled.
21	
22	In particular, two types of materials have found particular utility within the
23	present invention as being appropriate for efficient use in a separate layer,
24	inorganic bases and organic bases.
25	
26	Inorganic bases include metal oxides, hydroxides, and carbonates of Group IA
27	and like elements. Examples of suitable inorganic bases include calcium
28	carbonate, calcium hydroxide, potassium bicarbonate, and calcium oxide.

present invention.

1	
2	Organic bases can include any organic amine compounds including
3	amine-containing polymers, and preferably polyamine compounds. Organic
4	amine compounds are broadly defined as organic compounds containing at least
5	one amine group. Amine-containing polymers are defined as organic polymers
6	containing at least one amine group and includes grafting an amine group onto
7	the main polymer chain. Polyamine compounds are defined as any organic
8	polymer containing at least one amine group in each repeating unit. Preferably,
9	the organic amine compound is a non-volatile, non-migratory compound, e.g., it
10	does not migrate in the polymer system used.
11	·
12	Examples of suitable organic amine compounds include dipropylenetriamine;
13	tris(3-aminopropylene)amine, N,N,N'N'-tetrakis(3-aminopropyl)ethylenediamine
14	and 1,12 dodecanediamine. Examples of amine-containing polymers include
15	glycols containing amine groups such as polyethylene glycol with two amines
16	and polypropylene glycol with two amines, available from Texaco as Jeffamine;
17	and dimethylaminoethanol grafted ethylene-methyl acrylate copolymers.
18	Examples of polyamines include pentaethylene hexamine (PEHA); triethylene
19	tetraamine; polyvinyl oxazoline; and similar higher molecular weight compounds.
20	Such polymers can be used alone to form a film or can be reacted, blended, or
21	mixed with a film forming polymer.
22	
23	The neutralizing material is typically introduced into a thermoplastic resin such
24	as polyethylene, ethylene vinyl acetate, or ethylene methyl acrylate, in order to
25	improve its ease of processing. However, as long as it is in a form capable of
26	being introduced into the system and which form does not interfere with its
27	neutralizing functions, the method of introduction employed is not critical to the
28	present invention

PCT/US97/03307

1

The neutralizing materials are present in an amount which is effective to remove 2 3 at least a portion of the oxidation by-products or prevent such products from passing through. Preferably, it is present in an amount to significantly reduce or 4 eliminate migration of such by-products. 5

6

When employed with thermoplastic resins as discussed above, an effective 7 amount of the neutralizing material is typically from about 0.05-40% by weight 8 based on the second layer, depending on the particular neutralizing agent. For 9 10 example, where organic bases such as polyamine compounds are employed, the preferred amount is between about 0.05 and 40% by weight based on the 11 12 second layer.

13

14 The exact arrangement of the layers of the two components is not critical to the present invention as long as the oxidation by-products will come into contact 15 16 with the neutralizing material and thereby have their migration controlled.

17

18 19

20

It is preferred that the two layers be adjacent to each other. When used in a package, it is preferable that the neutralizing material be positioned between the contents of the package and the oxygen scavenging material, and therefore interim to the oxygen scavenging containing layer.

22 23

24

25

21

In one particular embodiment of the present invention, the two components are present in separate layers of a multi-layer film. The multi-layer structure of the present invention includes at least two layers:

26 27

a first layer comprising an oxygen scavenging material; and (a)

1	(b)	a second layer comprising at least one oxidation by-product neutralizing
2	,	material.
3		
4	The	multi-layer structure can include more than one of these neutralizing layers.
5		hough the arrangement of these layers is not critical to the present invention,
6		rder to improve its efficiency in preventing the migration of by-products into
7		interior of the package, as mentioned above, the second layer should be
8		ted between the first layer and the interior of the package.
9		
10	Pref	erably, this second layer is provided in conjunction with one or more
11	add	tional layers, e.g., oxygen barrier layers or polymeric selective barrier layers
12	that	also serve to aid in preventing the undesirable migration of the by-products
13	into	the package.
14		
15	Suci	n layers can include one or more of:
16		
17	(1)	An oxygen barrier layer which is typically an "outside" layer and comprises
18		a material which effectively functions as a physical barrier to oxygen, thus
19		minimizing or even eliminating any diffusion of oxygen into the resulting
20		package. The presence of such an oxygen barrier can supplement the
21		efforts of the neutralizing materials and therefore may reduce the amount
22		of neutralizing material needed.
23		
24		Although well recognized in the art, such layers typically have an OTR
25		(oxygen transmission rate) of no more than about 1 cc O ₂ /100 in. ² of
26		oxygen barrier layer/day/atm.
27		

1	(2	 A polymeric selective barrier layer as mentioned previously.
2	2	as mentioned previously.
3		For sake of completeness, the polymeric selective barrier layer functions
4		as a selective barrier to certain oxidation products but not to oxygen. In
5		one preferred embodiment, a layer is considered to be a polymeric
6		selective barrier layer when it prevents at least about half of the number
7 8		and/or amount of oxidation products having a boiling point of at least about
9		75°C from passing through the polymeric selective barrier layer from the
10		layer carrying the organic oxygen-scavenging material.
11		Polymoria calactica t
12		Polymeric selective barrier layers are typically located between the oxygen
13		scavenging layer and the "inside" of the resulting package to prevent
14		migration of the oxidation products into the package.
15		In fact, although the exact arrangement of these by-product blocking layers
16		is not critical to the present invention, it is preferred that both the second
17		neutralizing material layer(s), 4, and polymeric selective barrier layer(s), 3,
18		be located between the oxygen scavenging layer, 2, and the interior of the
19		package. Further, the oxygen barrier layer, 1, is typically located exterior
20		of the oxygen scavenging layer. Such an arrangement is illustrated in the
21		drawing figure.
22		
23	(3)	As further illustrated in the figure, the multi-layer film according to the
24		present invention can further include a sealing layer, 5, which is preferably
25 26		the innermost layer relative to the package formed therefrom. This layer
26 27		preferably comprises a heat sealable material.
41		

28

Optionally, where, for example, the second layer is the innermost layer, the heat 1 2 sealable material can be further included within the second layer containing the 3 neutralizing materials. 4 5 The multi-layer film according to the present invention can be produced by 6 conventional techniques, e.g., melt extrusion, co-extrusion, or lamination, which 7 are well recognized in the art. As such, they need not be described in detail 8 here. 9 10 Moreover, the multi-layer film according to the present invention can be 11 preferably employed in the production of packages, e.g., both rigid and flexible food packages, in the same manner as traditional multi-layer films. In a 12 13 preferred embodiment, the packaging and multi-layer film is UV transparent, in 14 order to allow sufficient UV radiation to be transmitted to the preferred oxygen 15 scavenging materials of the present invention. The preferred oxygen 16 scavenging materials are UV activated and therefore a UV transparent package 17 is critical to the system working. It is also preferred, particularly for food 18 packaging, that the package be optically clear. 19 20 The use of the neutralizing material in accordance with the present invention is 21 capable of removing a variety of the oxidation by-products produced by the 22 oxygen scavenging layer, particularly those odorous by-products. It is important, 23 however, that the neutralizing material of the present invention be kept in a layer 24 separate to that of the oxygen scavenging material in order to avoid inhibition of the oxidation reaction. The efficiency of the present invention in removing these 25 26 by-products is particularly apparent when employed in combination with the 27 polymeric selective barrier layer in a multi-layer film.

1	The oxygen scavenging system of the present invention can find broad utility
2	with a variety of oxygen sensitive materials. Such materials include, in addition
3	to food, cosmetics and beauty products, other oxygen sensitive chemicals and
4	electronic materials.
5	
6	For example, in a packaging environment, in addition to being present as a film
7	which forms at least a portion of the package, it can be found in virtually any part
8	of a packaging material which will come into contact with the interior of the
9	package. Such environments include but are not limited to cap liners for bottles
10	and the like, trays, e.g., those trays used in the food industry, packaging
11	materials for cosmetic or other materials in the beauty industry, as well as other
12	chemical environments.
13	
14	The present invention will now be discussed in terms of certain examples, it
15	being understood that such examples are solely illustrative in nature and in no
16	way limit the present invention.
17	
18	EXAMPLES
19	·
20	Examples 1-4
21	_
22	Examples 1-4 are of three layer structures having a A/B/A arrangement at 1:1:1
23	thickness with a total thickness of 3 mils. In each case, A is ethylene methyl
24	acrylate copolymer, EMAC® SP-2260, with the named additive, and B is
25	ethylene methyl acrylate benzyl acrylate copolymer containing 1000 ppm of
26	cobalt in the form of cobalt neodecanoate.

Example	Additive	Odor
1 Control	None	acetic
2	2% PEHA polyamine	weak acetic with amine smell
3	0.5% PEHA polyamine	
4	2% calcium carbonate	weaker acetic

 All films were irradiated for 5 minutes under a UV blackray (250 nm) at a 5-inch distance. The oxygen scavenging resins were sealed in one-liter size bottles and oxygen uptakes were monitored by Mocon 710 oxygen meter for 30-40 days using 2 gram samples. At the end of the oxygen uptake of about 100 cc/gram resins, the bottles were opened and odor evaluated by a panel of at least 3 people. The results are set forth in the foregoing table.

Based upon the foregoing results, it can be concluded that a small amount of neutralizing agent in a separate layer is able to significantly reduce undesirable acetic smell after the products were extensively oxidized. Since a polymer selective barrier can be inserted and real application conditions involve reduced oxygen (<2% O₂ after nitrogen flush compared to 20% in the present examples), the present invention is expected to significantly reduce by-product odor.

1		Example 5				
2		<u>Example o</u>				
3	Αn	A multi-layer film construction was made having from the outer most layer to the				
4	inn	inner most layer (i.e., that to be generally adjacent to foods) the following layers				
- 5						
6	(1)	a 0.16 mil thick aluminum foil oxygen barrier layer;				
7						
8	(2)	a 1 mil thick EBZA oxygen scavenging polymer with 750 ppm of Cobalt in				
9		the form of cobalt neodecanoate salt;				
10						
11	(3)	a 0.5 mil thick oriented PET functional barrier layer; and				
12	(4)					
13	(4)	a 1.0 mil thick EMAC 2205 containing 2% of calcium carbonate as the				
14 15		neutralization layer as well as heat seal layer.				
16	The	films word IN Granding at a				
17	2000	films were UV irradiated from the inside layer out as described in				
18	piece	rdance with Examples 1-4 and 9" by 9" bags are made by heat sealing two				
19	conta	es of the multilayer construction together. The bags were then inflated to				
20	Moco	contain 1 liter fresh air by a syringe. The oxygen content was monitored by a Mocon 710 oxygen meter.				
21		and any gent mater.				
22	At the	e end of 4 weeks, 80% of the oxygen was consumed and the bag was				
23	open	ed slowly for odor evaluation as described in accordance with				
24	Exam	iples 1-6.				
25						
26	A con	strol construction containing identical layers, except no calcium carbonate				
27	was u	ised in the EMAC layer, was also tested. The control gave a strong acidic				
28	smell.	The opened bag of the present invention gave no detectable acidic odor.				
		5 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 -				

2	This example further demonstrates the effectiveness of the present invention in
3	controlling odor even though the neutralizer layer is separate from the oxygen
4	scavenging polymer.
5	
6	Example 6
7	
8	This example demonstrates the use of an amine-containing polymer in odor
9	removal. Ester exchange of EMAC with N,N-dimethylethanolamine was
10	conducted in a twin screw extruder by reacting 4.5 Kg/hr EMAC® SP 2260
11	containing 24% by weight methylacrylate and having a melt index of 2,
12	1.13 Kg/hr N,N-dimethylethanolamine, and 45 cc/hr titanium tetraethoxide at
13	210°C. The conversion was 22.1% to N,N-dimethylethyl ester of acrylate.
14	and the right difficulty letter of acrylate.
15	A two-layer film structure with a total thickness of 4-5 mil was prepared by
16	coextrusion of the above amine-containing polymer and EBZA (47.2%
17	conversion) containing 1000 ppm cobalt, in the form of cobalt neodeconate. The
18	two-layer film was aged in a one-liter bag (21% oxygen) at room temperature for
19	about one month. The bag was opened and the odor compared by a three-judge
20	panel. The odor was slightly less acidic than the monolayer film without the
21	amine-containing polymer.
22	
23	Principles, preferred embodiments, and modes of operation of the present
24	invention have been described by the foregoing. The invention is not to be
25	limited by particular embodiments disclosed since they are only illustrative in
26	scope.
27	

- Various modifications, substitutions, omissions, and the like, may be made 1
- without departing from the spirit of the invention. Accordingly, it is intended that 2 3
- the scope of the present invention be limited solely by the scope of the following 4
- claims including equivalents thereof.

1	WH	HAT IS CLAIMED IS:				
2						
3	1.	A multi-layer structure useful in the packaging of an oxygen sensitive				
4		material, the structure comprising:				
5		-				
6		(a) a first layer comprised of an oxygen scavenging material which forms				
7		at least one by-product upon the reaction thereof with oxygen; and				
8						
9		(b) a second layer separate from the first comprised of an effective				
10		amount of neutralizing material capable of neutralizing by-products				
11		formed upon the reaction of an oxygen scavenging material with				
12		oxygen.				
13						
14	2.	The multi-layer structure of claim 1, wherein the structure is UV transparent				
15		and optically clear.				
16						
17	3.	The multi-layer structure of claim 1, wherein the structure has more than				
18		two layers.				
19						
20	4.	The multi-layer structure of claim 1, wherein the structure further comprises				
21		an oxygen barrier layer.				
22						
23	5 .	The multi-layer structure of claim 1, wherein the structure further comprises				
24		a selective barrier layer.				
25						
26	6.	The multi-layer structure of claim 1, wherein the second layer is interior to				
27		the first layer.				

?	The multi-layer structure of claim 1, wherein the oxygen scavenging material comprises an organic oxidizable material.
	The multi-layer structure of claim 1, wherein the oxygen scavenging material comprises a polymeric material having oxidizable sites.
9.	The multi-layer structure of claim 8, wherein the oxygen scavenging material further comprises a transition-metal salt catalyst.
10.	The multi-layer structure of claim 9, wherein the transition-metal salt catalyst is cobalt (II) neodecanoate, cobalt (II) oleate, cobalt (II) linoleate, or cobalt (II) caprylate.
11.	The multi-layer structure of claim 1, wherein the neutralizing material is selected from the group consisting of inorganic bases and organic bases.
12.	The multi-layer structure of claim 11, wherein the organic base comprises an organic amine compound.
13.	The multi-layer structure of claim 1, wherein the second layer further comprises a thermoplastic resin.
14.	The multi-layer structure of claim 1, wherein the neutralizing material in the second layer comprises an acid-, alcohol- or aldehyde-reactive material.
15.	The multi-layer structure of claim 14, wherein the neutralizing material comprises an inorganic base or an organic amine compound.
	10. 11. 12. 13.

1	16	The multi-layer structure of claim 15, wherein the neutralizing material is
2		selected from the group consisting of calcium oxide, calcium hydroxide,
3		potassium bicarbonate and calcium carbonate.
4		
5	17.	The multi-layer structure of claim 15, wherein the organic amine comprises
6		an amine-containing polymer.
7	•	
8	18.	The multi-layer structure of claim 15, wherein the organic amine compound
9		comprises a polyamine compound.
10		
11	19.	The multi-layer structure of claim 1, wherein the structure further comprises
12		an oxygen barrier layer which is located outside of the first layer and a
13		polymeric selective barrier layer which is located interior to the first layer.
14		
15	20.	The multi-layer structure of claim 19, wherein the structure further
16		comprises a seal layer which is the innermost layer of the structure.
17		,
18	21.	The multi-layer structure of claim 19, wherein the oxygen scavenging
19		material is selected from the group consisting of organic oxidizable
20		materials and polymeric materials having oxidizable sites, and the
21		neutralizing material is selected from the group consisting of inorganic
22		bases and organic bases.
23		
24	22.	The multi-layer structure of claim 19, wherein the organic base comprises a
25		polyamine compound.
26		

	2.	or the multi-layer structure of claim 22, wherein the polyamine compound is
2	2	selected from the group consisting of pentaethylene hexamine, triethylene
3	3	tetraamine, and polyvinyl oxazoline.
. 4		
5	24	The multi-layer structure of claim 19, wherein the inorganic base is
6		selected from the group consisting of calcium oxide, calcium hydroxide,
7		and calcium carbonate.
8		
9	25	A package suitable for holding an oxygen scavenging material comprising
10		a multi-layer structure, wherein the multi-layer structure is that structure
11		according to claim 1.
12		
13	2 6.	The package according to claim 25, wherein the structure further
14		comprises an oxygen barrier layer which is located outside of the first layer.
15		· · · · · · · · · · · · · · · · · · ·
16	27.	The package according to claim 25, wherein the structure further
17		comprises a polymeric selective barrier which is located inside of the first
18		layer.
19		
20	28.	The package according to claim 25, further comprising a polymeric
21		selective barrier layer which is between the first and the second layer.
22		
23	29 .	The package according to claim 25, further comprising at least one of a
24		chemical material, or an electronic component located therein.
25		
26	3 0.	The package according to claim 25, further comprising a food product or
27		cosmetic product located herein.
28		

1	31.	The package according to claim 25, wherein the multi-layer structure is a
2		film which forms at least a portion of the package.
3		
4	3 2.	The package according to claim 25, wherein the multi-layer structure is a
5		film which is present within the package.
6		
7	33 .	The package according to claim 25, wherein the package, together with a
8		multi-layer structure, is optically clear and UV transparent.
9		
10	34 .	An article for packaging an oxygen sensitive product comprising an oxygen
11		barrier and a non-integral packaging component selected from the group
12		consisting of coatings, bottle cap liners, adhesive or non-adhesive inserts,
13		sealants, gaskets and fibrous mat inserts, wherein the non-integral
14		packaging component comprises an oxygen scavenger and a by-product
15		absorber.

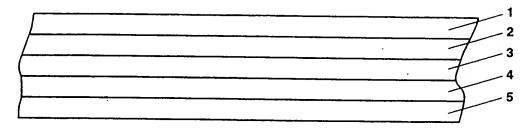
	**************************************	A PROPERTY AND A STATE OF THE S	হৈ ্যুক্ত হ	्रा वर्षे जिल्लाम् <mark>य</mark>	
:					
			/1.2.		

		*			
	* }; 	,			e and the second
			8		
•	.*.				
					•
4	_		*	*	
· - <u>-</u>	ンデ			· -	* 8
				*	→
•		•			
				2	
•			-		. 7
· ·		, , , , , , , , , , , , , , , , , , ,	A contract of	•	• - ()
* * * * * * * * * * * * * * * * * * * *					
		*			
•		a de la companya de l	· .	•	. ***
•					•
****		**		* • •	
* * * * * * * * * * * * * * * * * * * *		· · · · · · · · · · · · · · · · · · ·			
		0 0 0			
			3. ·	& t	
		* * *			*
	*				
				Nag* .	. *
					. * >
· 16 %					
			*		. *
					*
					*
	*				
	*				

1/1

FIGURE 1

Exterior



Interior

į.				
	•			
1		. *		
r				
1				e (leg
	. •		*	
R.				
Se.		•		
			· · · · · · · · · · · · · · · · · · ·	
		,	• .	
			•	
File July	• .			
		1	* * * * * * * * * * * * * * * * * * * *	
· ·				
ž.				
	•			
			$\eta_{ij} = \eta_{ij}$	
			*	
			*	
St.			×	
E	. 20			
-				
	* * *			
. II.	¥ 2			
A ⁵	· · · · · · · · · · · · · · · · · · ·		• •	
*	•			
;.			A. A.	
			e e e	
	* .			
4.				
			•	
			10	
			•	(f)
	je "			
	*			
Ť				
			(a) 4	
				*
	. * *			
	· ·		. •	•
-	v			
			, es	
	• •			
•	*			
· · ·	•			
•	•			
i⊬ ∡ji	•		25	
	* ×		**	
	· · ·			
•			-2.4	
	*			
No.				
_		. 76 % 4.		1 (Garage 25) 1 () 1 () () () () () ()

INTEL: ATIONAL SEARCH REPORT

Inter onal Application No PCT/US 97/03307

A. CLASSIFICATION OF SUBJECT MATTER IPC 6 B32827/18 B65D81/26 A23L3/34 According to International Patent Classification (IPC) or to both national classification and IPC **B. FIELDS SEARCHED** Minimum documentation searched (classification system followed by classification symbols) B32B B65D IPC 6 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) C. DOCUMENTS CONSIDERED TO BE RELEVANT Relevant to claim No. Citation of document, with indication, where appropriate, of the relevant passages Category * 1,3,4,6, EP 0 638 487 A (TOYO SEIKAN KAISHA LTD) 15 X 13,25, February 1995 26,31,32 see page 2, line 5 - line 9; claims; figures 3,4 see page 3, line 22 - page 5, line 8 see page 7, line 51 - page 8, line 19 2,5, Y 7-12, 14-24, 27-30, 33,34 -/--Patent family members are listed in annex. Further documents are listed in the continuation of box C. Х Special categories of cited documents: "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the document defining the general state of the art which is not considered to be of particular relevance invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone earlier document but published on or after the international filing date document which may throw doubts on priority claim(s) or 'Y' document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person stilled in the art. which is cited to establish the publication date of another citation or other special reason (as specified) document referring to an oral disclosure, use, exhibition or other means document published prior to the international filing date but later than the priority date claimed "&" document member of the same patent family Date of mailing of the international search report Date of the actual completion of the international search O5, O4 97 22 July 1997 Authorized officer Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentiaan 2 NL · 2280 HV Rijswijh Tel. (+ 31-70) 340-2040, Tx. 31 651 epo nl. Pamies Olle, S Fax: (+31-70) 340-3016

INTERNATA NAL SEARCH REPORT

Intr onal Application No PCT/US 97/03307

Category *	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	PCT/US 97/03307
85.7	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 93 01049 A (FRITO LAY INC; DU PONT (US)) 21 January 1993 cited in the application see page 3, paragraph 3 - page 4, paragraph 1; claims; table 1 see page 5, last paragraph - page 6, paragraph 1	11,12, 14-18, 20-24
Y	US 5 211 875 A (SPEER DREW V ET AL) 18 May 1993 cited in the application see column 3, line 36 - line 39; claims see column 4, line 44 - column 5, line 12 see column 7, line 58 - column 8, line 60	2,5, 7-10,19, 21, 27-30, 33,34
A	DATABASE WPI Section Ch, Week 9519 Derwent Publications Ltd., London, GB; Class A92, AN 95-143805 XP002035800 & JP 07 067 594 A (TOYO SEIKAN KK), 14 March 1995 see abstract	1,3,4, 25,30-32
	EP 0 464 782 A (VISKASE CORP) 8 January 1992 cited in the application see page 2, line 3 - line 6; claims 1,8,10,15 see page 2, line 29 - line 44	1
.A .	EP 0 720 821 A (SUMITOMO CHEMICAL CO) 10 July 1996 see page 2, line 48 - line 50; claims 1,4,10,18,19 see page 4, line 9 - line 11 see page 4, line 39 - line 52	1,3
TASAGIO	stinuation of second sheet) (July 1992)	